

28(3) S/028/60/000/01/015/033 D041/D002 AUTHOR: Zembovskiy, I.F., and Filippov, M.M.

Unification of Parts by Using the Group TITLE:

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of Machining

PERIODICAL: Standartizatsiya, 1960, Nr 1, pp 42-43

ABSTRACT: Some machine building plants, and particularly

the Laptevskiy zavod ugol'nogo mashinostroyeniya (Laptevo Coal Mining Machine Plant) have started using the "group method" for machining parts, suggested by S.P. Mitrofanov, Lenin prize laureate. The method consists in splitting machine parts into groups of similar configuration, dimensions, and according to the required production processes. For every group of parts, special machining equipment is produced, and machine tool attachments adjusted. The "group method" considerably reduces the quantity of equipment required, cuts expenses, and increases the work productivity by 25 to 40%.

Card 1/1

25(5)

SOV/28-59-2-9/26

AUTHORS:

Zembovskiy, I.F. and Filippov, M.M., Engineers

TITLE:

Unification of Rubber Sealings (Unifikatsiya uplotneniy iz

reziny)

PERIODICAL:

Standartizatsiya, 1959, Nr 2, pp 32-33 (USSR)

ABSTRACT:

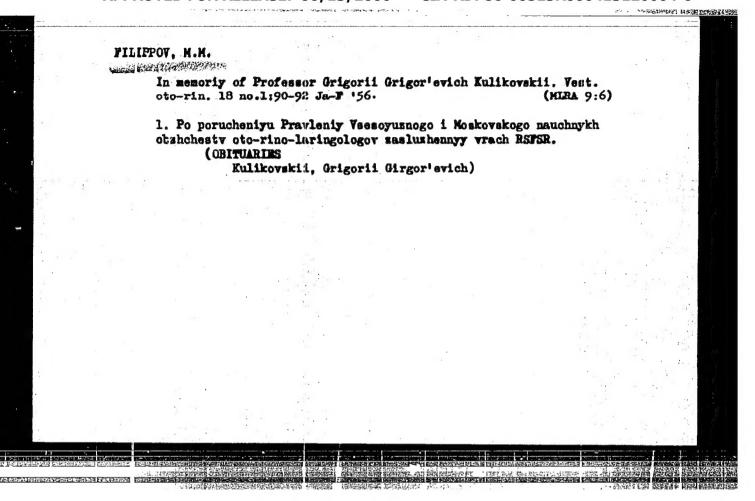
The Laptevo Coal Mining Machine Building Plant uses many different rubber sealings and components with varying rubber bases. The plant's office for standardization and normalization reduced the number of rubber grades from 11 to 4 without affecting the quality of the product. The authors stress the need to standardize manufacture of circular section rubber sealing rings. At present the Tula and Moscow technical rubber equipment plants manufacture the rings from press-forms designed and produced by the plants themselves. Centralized production of these rings by specialized plants will cut down the production costs.

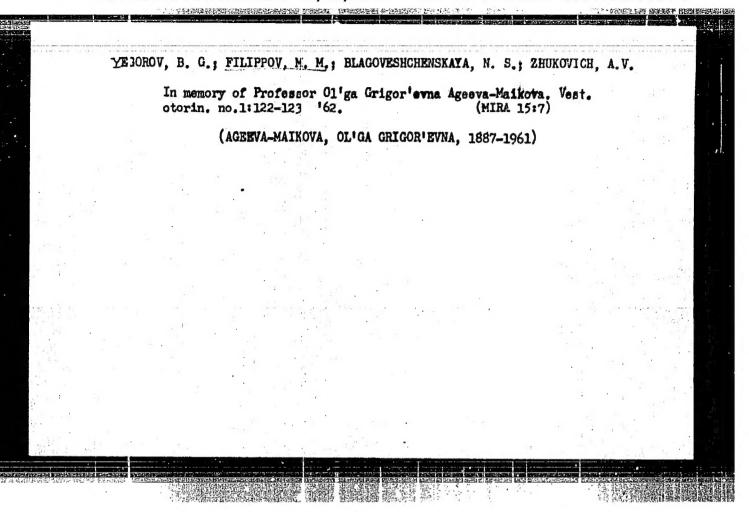
ASSOCIATION:

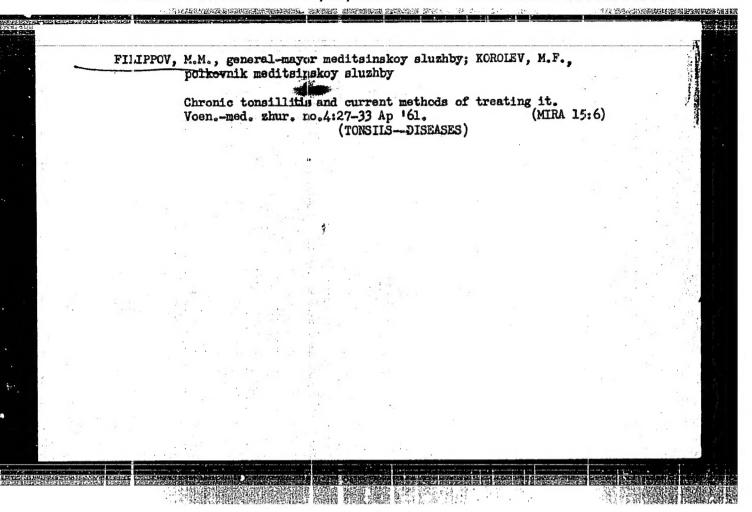
Laptevskiy zavod "Uglemash" (The Laptevo "Uglemash" Plant)

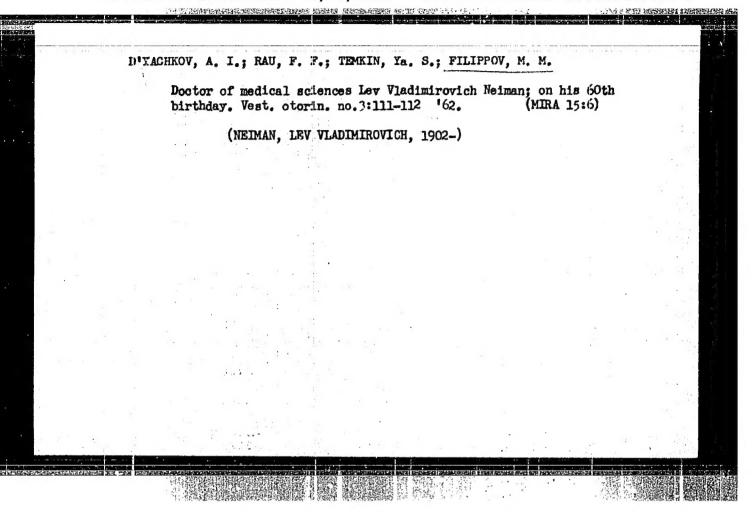
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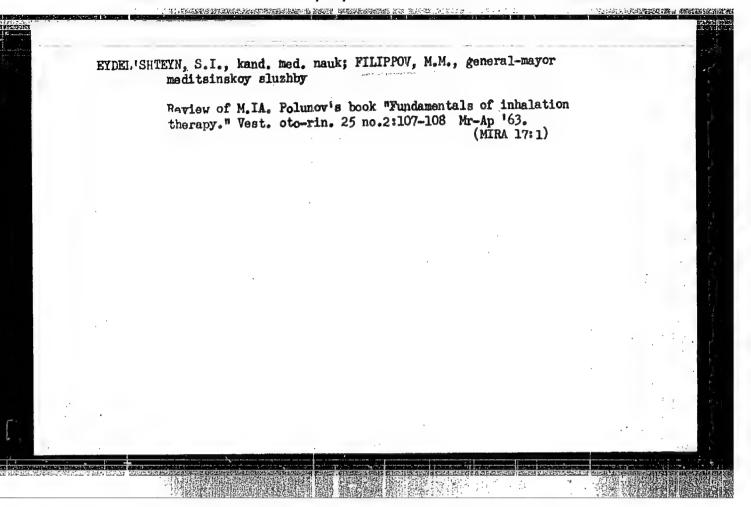
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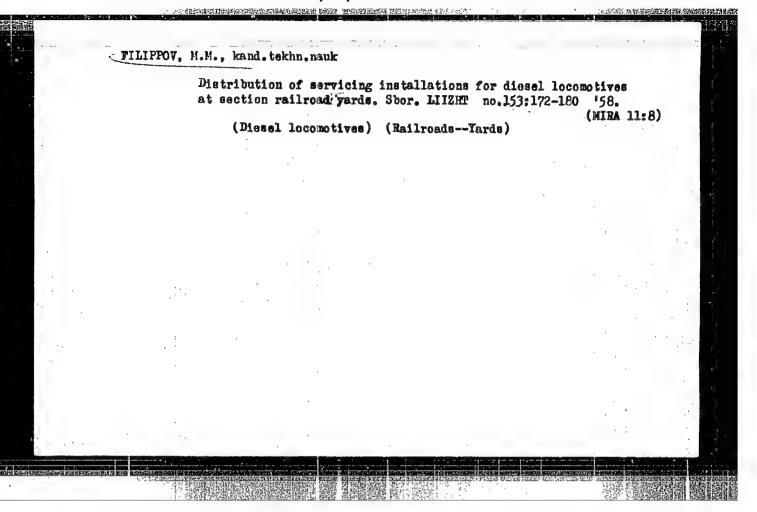


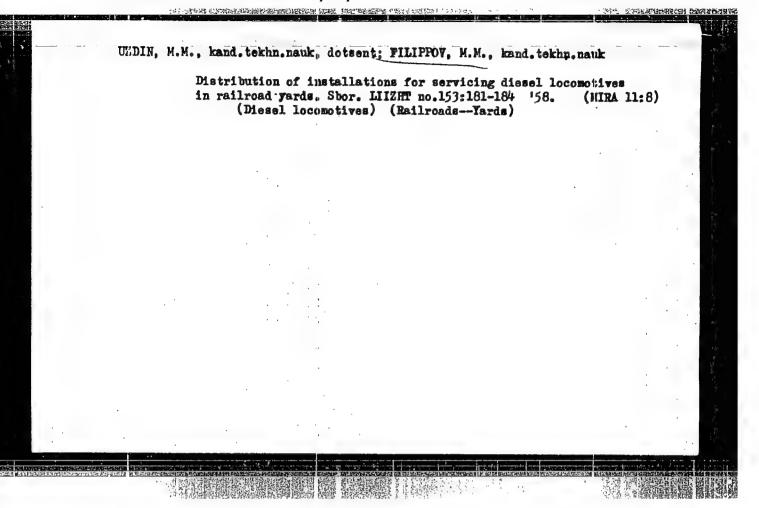


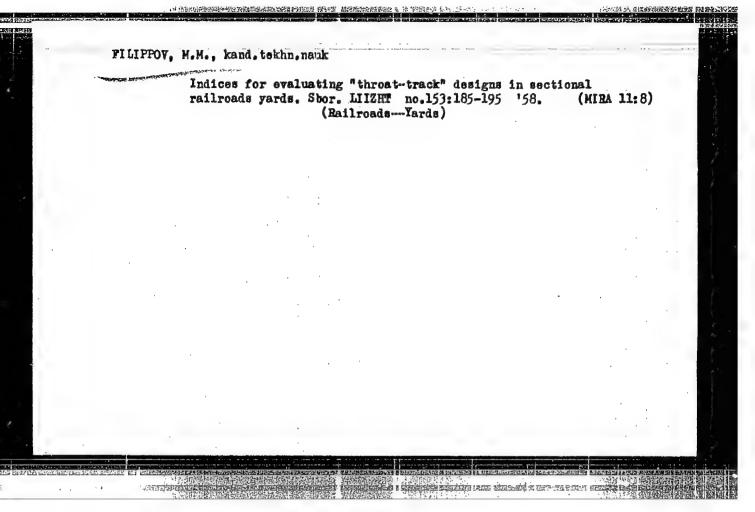


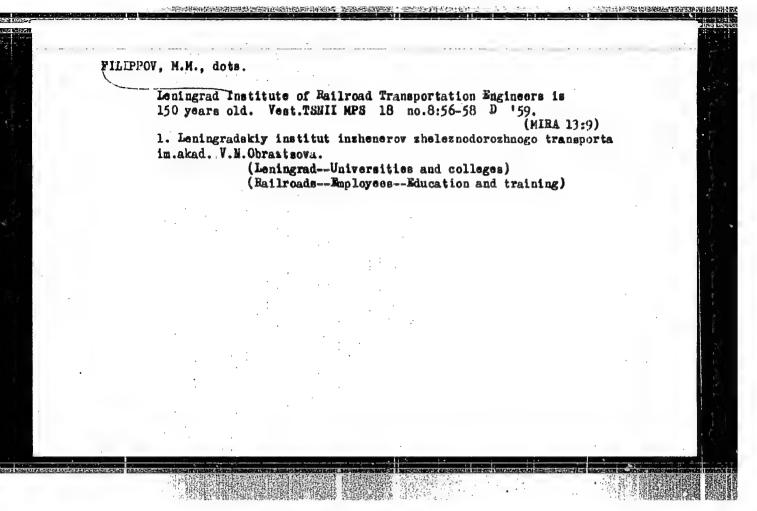








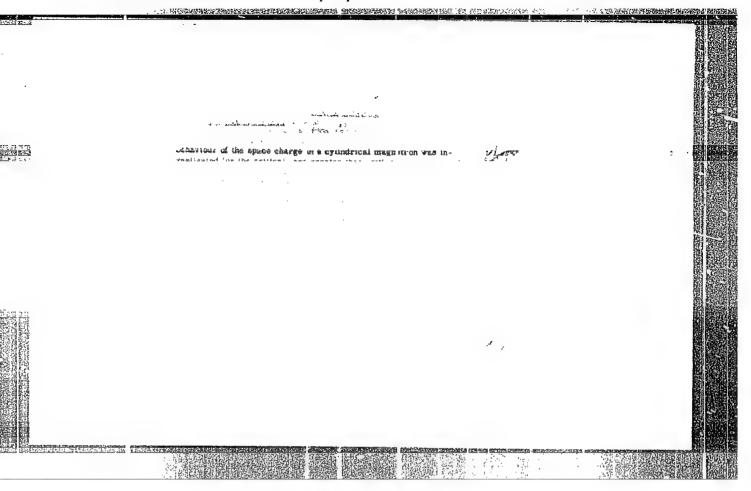




M. M. FILIPPOY, MOLCHANOV, A. P., E. M. GYUNNINEN, A. V. MEL'HIKOV, AL. P. MOLCHANOV, L. MYASNIKOV, V. N. RYSAKOV, F. I. SKRIPOV

"Results of Solar Eclipse Observations of 1952 and 1954 in the 3.2 cm Wavelength"

(Total Eclipse of the Sun, February 25, 1952 and June 30, 1954, Transactions of the Expedition to Observe Solar Eclipses) Moscow, Izd-vo AN SSSR, 1958. 357 p.



1 i PROV. M. M. 120-3-22/40

AUTHORS: Rakovskiy, I.I. and Filippov, M.M.

TITLE: A Broad-Band Oscillator Using a Lighthouse Triode. (Shirokodiapazonnyy generator na mayachkovom triode)

PERIODICAL: Pribory i Tekhnika Eksperimenta, 1957, Nr 3, pp.80-81 (USSR)

ABSTRACT: A decimeter-band oscillator using a lighthouse triode 6C5A is described. The oscillator can be tuned over the 12-60 cm band with sufficient power output for experimental work. The arrangement is shown in Fig.1. The two-conductor line consists of two lengths of copper wire 3 mm dia and 20-30 cm (or more) long, connected to the anode and grid discs respectively. A capacitor C1 (tens of pFs) is connected by sliding contacts across the lines and its posi-

connected by sliding contacts across the lines and its position can be changed to suit a particular frequency. The remaining part of the line has similar capacitors across it, separated from each other by  $\lambda/2$ . The metallic base of the tube is completely enclosed in a brass cylinder, 1, the height of which is several mm greater than the height of the base. Between this cylinder and the grid disc is connected a loop, 2, made of thick foil or wire, which makes DC contact between the grid and the cathode. The distributed capacity of this loop and the distributed capacity and

Card 1/2

120-3-22/40

A Broad-Band Oscillator Using a Lighthouse Triode.

inductance of the cylinder form the grid resonant circuit. Connection of the loop also alters the inter-electrode capacity forming the feedback path. Altering the dimensions of the loop alters the frequency of oscillation. The energy is taken off by a coupling element or by an antenna. By selecting the second harmonic waves down to  $\lambda = 12$  cm can be obtained. There is I figure and no references.

ASSOCIATION: Leningrad State University im. A.A.Zhdanov. (Leningradskiy gosudarstvennyy universitet im. A.A.Zhdanova)

SUBMITTED: January 16, 1957.

AVAILABLE: Library of Congress.

Card 2/2 1. Oscillators-Operation 2. Triode-Applications

PILIPPOV, M:M.; BUKIN, A.N.

Oscillograph for the centimeter band. Ixv. vys. ucheb. zav.; radiotekh. no.3:373-376 My-Je '58. (MIRA 11:7)

1.Rekomendevana kafedroy radiofiziki Leningradskogo gosudarstvennege universiteta.

(Oscillograph) (Microwaves)

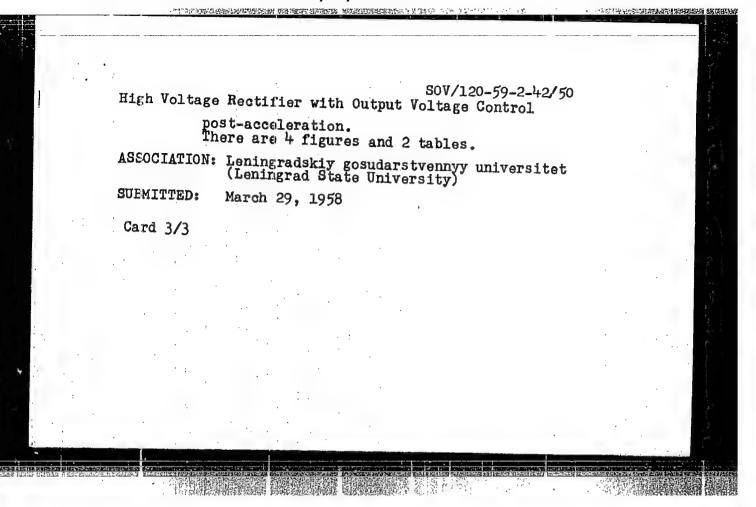
SOV/120-59-2-42/50 AUTHORS: Bukin, A.N., and Filippov, M.M. TITLE: High Voltage Rectifier with Output Voltage Control (Vysokovolitnyy vypryamitel' s reguliruyemym vykhodnym napryazheniyem) PERIODICAL: Pribory i tekhnika eksperimenta, 1959, Nr 2, pp 139-141 (USSR) ABSTRACT: The block diagram is shown in Fig 1. The low voltage supply, which should be stabilised, is first converted into a voltage between 12 and 15 kV peak at a frequency of 8 kc/s. This derived supply is rectified in 3 units. In the first two units D.C. supplies of +10 kV and -10 kV are formed. The third unit is a voltage quadrupler with outputs at +20, +30 and +50 kV. There are auxiliary units for feeding the heaters of the high voltage rectifiers in the quadrupler period. Fig 2 is a more detailed circuit diagram, and Table 1 describes the five-winding coil to which the GU-29 valve is connected as a Hartley oscillator. Fig 3 shows how the coil is mounted with respect to the first two rectifiers. Card The oscillator is tuned by varying the position of a cylindrical ferrite core of F-600 material and its 1/3

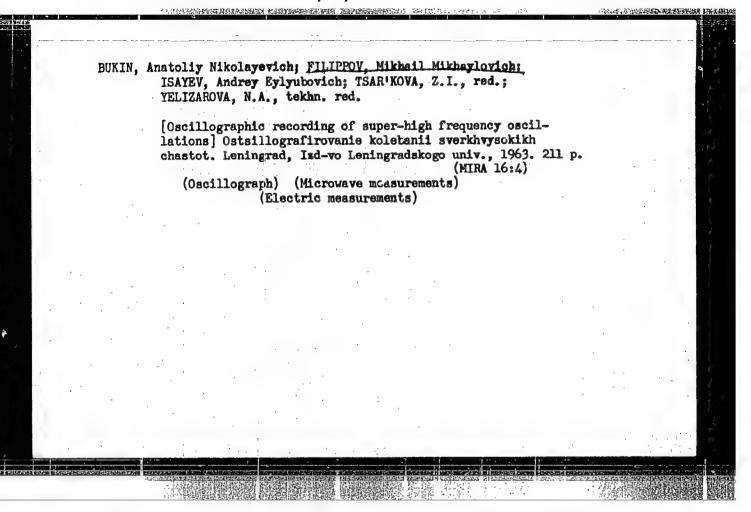
SOV/120-59-2-42/50

High Voltage Rectifier with Output Voltage Control

output voltage can be varied from 2 to 12 kV by changing the grid bias but this results in a dissimilar variation of the ± 10 kV outputs. The quadrupler uses valves type 1Ts11P and would normally supply +40 kV for an input of 11 kV. By connecting the rectifier system in series with the +10 kV supply a maximum of +50 kV can be obtained. The +20 kV supply is filtered via the components R4C12. Valves  $\mathcal{N}_8$  and  $\mathcal{N}_9$  are a pulse power supply for the quadrupler heaters and are coupled by the transformer TI-1 shown in Fig 4. Table 2 gives some typical readings taken at various points in the circuit. The quadrupler together with its heater transformer is mounted separately in a unit measuring 130 x 170 x 110 mm3 completely filled with paraffin. The whole arrangement measures 300 x 300 x 200 mm3 and can supply 1 mA at 50 kV to a UHF oscillograph tube using

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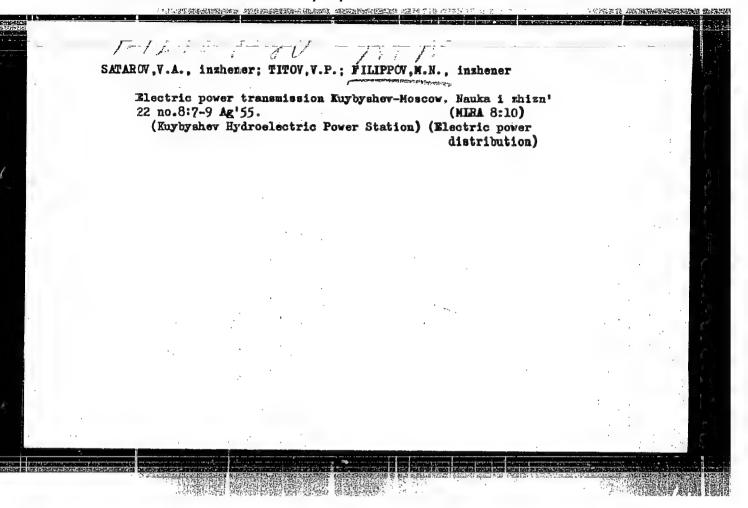


FILIPPOV. M. II.; PLOSHCHANIKOVA, YE. A.

Jerusalme Artichoke

Cultivating the Jerusalem artichoke. Korm. baza 3 no. 7, 1952.

Monthly List of Russian Accessions, Library of Congress, September 1952. UNCLASSIFIED.



22(1)

SOV/3-59-3-3/48

AUTHOR:

Filippov, M.P.

TITLE:

A Broad Road to the Vuzes for Production Workers (ProizVodstvennikam - shirokuyu dorogu v vuzy)

PERIODICAL:

Vestnik vysshey shkoly, 1959, Nr 3, pp 8-11 (USSR)

ABSTRACT:

The author tells of the experience gained by the Leningrad vuzes in training production workers. In 1957, the number of production workers enrolled in the vuzes amounted to 5,569, i.e. 30.2% of the total number; in 1958, it was 7,681 with almost half of the freshmen having a 2-year record of practical work. In 4 vuzes of the city - the Sel'skokhozyaystvennyy, Veterinarnyy, Bibliotechnyy institut (Agricultural, Veterinary, Library Institute) and the Institut sovetskoy torgovli (Institute of Soviet Trade) - 80% of the freshmen were production workers. Many students have interrupted their education for a considerable time, in some cases for 8 to 10 years. The City Committee of the Party advised the Party organizations of vuzes to

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A Broad Road to the Vuzes for Production Workers

especially help these students so that they won't drop out in the first term. Poor students were granted scholarships, and those from other towns were accomodated in hostels. To overcome the educational gaps, many institutes organized additional studies in mathematics, physics, chemistry, etc. for the production workers. At present the level of training of those admitted in the fall of 1958 compares favorably with the production workers who were accepted in 1957. An analysis of last year's results made in 6 institutes - the Polytechnical, Electrical Engineering imeni V.I. Ul'yanov (Lenin), Pedagogical imeni Gertsen, Engineering and Construction, 1st Medical, and Mechanical - leads to the conclusion that the majority of the production workers were able to cope with the training program. In several subjects they proved to have even a better knowledge than those who had come direct from school. The author comments on these results giving numerical data in respect to the individual institutes. The Leningrad

Card 2/3

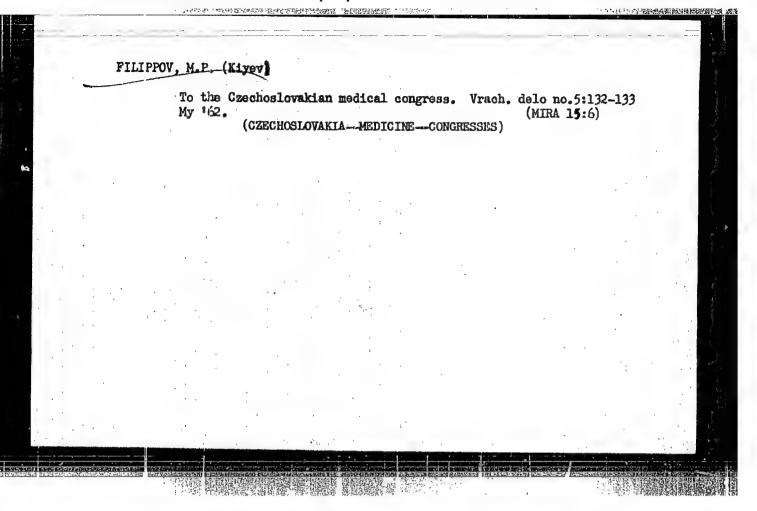
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A Broad Road to the Vuzes for Production Workers

KPSS City Committee has recommended that a pre-admission training of the students be developed. This year the short-term courses are planned to last 9 months.

ASSOCIATION: Leningradskiy gorodskoy komitet KPSS (Leningrad City Committee of the KPSS)

Card 3/3



#### "APPROVED FOR RELEASE: 06/13/2000

CIA-RDP86-00513R000413120004-6

M. P. FILIPPOV USSR/Physical Chemistry - Molecule, Chemical Bond. B-4 Abs Jour : Referat Zhur - Khimiya, No 1, 1958, 99 Author A.V. Ablov, M.P. Filippov. Inst Title : Light Absorption by Complex Compounds of Trivalent Cobalt. I. Dependence of Absorption Spectra of Compounds of Co-En\_Amine Cl\_/X, Type on Nature of Co-ordinated Amine. Orig Pub : Zh. neor@an. khimii, 1957, 2, No 1, 42-52 Abstract : The absorption spectra of solutions of cations./CoEn\_Amine Cl/ in 0.1 n. HCl were investigated. Ethylamine, benzylamine, aniline, n-fluoraniline, n-chloraniline, m-toluidine, o-anisidine, O-phenetidine and n-anisidine were the amines in this cations. 3 absorption bands were found in the absorption spectrum of the cations  $(CoEn_2(XC_6H_1NH_2)C1)^{2+}$  at 520, 305 to 345 and 232 to 235 m. A 4th absorption band was also observed at 215 to 218 mm in some cases (amine = n-chloraniline, 0-anisidine and o-phenetidine). Card 1/3 Richiner State Univ

USSR/Physical Chemistry - Molecule, Chemical Bond.

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Abs Jour : Ref Z

: Ref Zhur - Khimiya, No 1, 1958, 99

The 1st and 3rd absorption bands little depending on the nature and position of the substitute in the aniline nucleus are attributed to the groupation En<sub>2</sub>CoCl(C<sub>6</sub>H<sub>5</sub>NH<sub>2</sub>). The stability of the position of the 1st band notwithstanding the differences in the limits of the fluctuations of dipole moments of the amines (from 1.51 to 2.97) contradicts the electrostatic theory of Hartmenn (RZhKhim, 1956, 3162). The position of the 2nd band depends on the nature of the substitute, as well on the substitution type, and its bathochromic shift rises in the series n-F, m-CH<sub>3</sub>, n-Cl, n-CH<sub>3</sub>, 0-OCH<sub>3</sub>, o-OC<sub>2</sub>H<sub>5</sub>, and n-OCH<sub>3</sub>. The

absorption spectra of \[ \tilde{\text{CoEn}}\_2(\text{NH}\_2\text{C}\_2\text{H}\_5)\text{Cl}\]^{2+} and \[ \tilde{\text{CoEn}}\_2-\text{(NH}\_2\text{CH}\_2\text{C}\_6\text{H}\_5)\text{Cl}\]^{2+} are very close and differ only in the

left hand branch of the short wave band. Considering

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USSR/Physical Chemistry - Molecule, Chemical Bond.

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Abs Jour

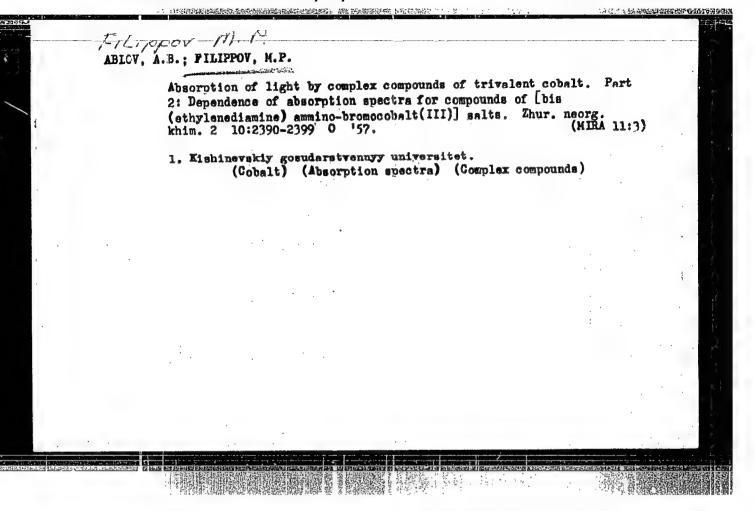
: Ref Zhur - Khimiya, No 1, 1958, 99

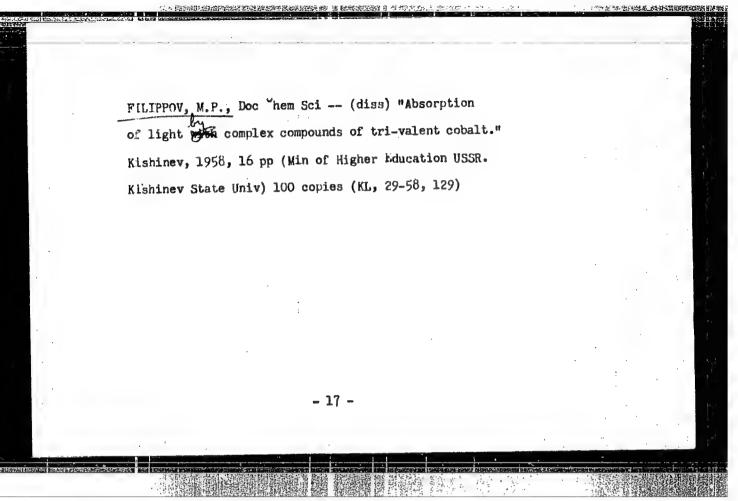
the similarity of the absorption spectra of these two cations with the spectrum of 1.2-/CoEn<sub>2</sub>NH<sub>3</sub>Cl/<sup>2+</sup>, the

authors arrive at the conclusion regarding the cis-structure of the investigated cations. It is postulated that the structure of all the other studied cations is also a cis-structure. The synthesis of CoEn2.(n-C6H4NH2C1)C1/C12.

.H<sub>2</sub>0 and  $\sqrt{\text{CoEn}}_2(\text{NH}_2\text{C}_2\text{H}_5)\text{Cl}/\text{S}_2\text{O}_6.\text{H}_2\text{O}$  is described.

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AUTEORS: Ablov, A.V., Filippov, M.P. SOV/78-3-7-17/44 III. The Dependence of the Absorption Spectrum of the Compound TITLE: of the Type Co.Amin<sub>2</sub> (DH) 2 x on the Nature of the Coordination (III. Zavisimost' spektrow pogloshchemiya soyedineniy tipa Co.Amin<sub>2</sub> (DH)<sub>2</sub> ot prirody koordinirovannogo amina) PERIODICAL: Zhurnal neorganicheskoy khimii, 1958, Vol. 3, Nr 7, pp. 1565-1572 (USSR) ABSTRACT: The absorption spectrum within the range of A = 210-650 m A up to dimethylglyoxymodiaminecobalt-cation complex [Co (Amin2 (DH)2]+ was investigated, in which amine = ammonia, pyridine, aniline, o., m. and p-toluiding, m- and p-ohloroaniline, m- and p-bromoaniline, p-iodine aniline, o- and p-anisidine and o-phenetidine. DH demotes the remainder of dimethyl glyoxin. The first line of the investigated compounds shifts in the direction of the short wave range in contrast to the corresponding ethylene-diamine compounds. The occurrence of new intense lines in the absorption spectrum on the introduction of aniline into the inner sphere of the complex is caused by the group  $\text{Co-N} \longrightarrow$ . During exchange into the complex  $\left[\text{Co (NH}_3)_2 \left(\text{DH}\right)_2\right]^+$  of one molecule ammonia for Card 1/2

III. The Dependence of the Absorption Spectrum of the Compound of the Type [Co.Amin2 (DH) 2 x on the Nature of the Coordination Amine

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SOV/78-3-7-17/44

ome chlorine atom the position of the lines of the absorption spectrum changes only slightly. The occurrence of the lines in the absorption spectrum at 250 m  $\mu$  does not depend on the nature of the amine but upon the groups.

$$CH_3 - C = N$$

$$CO = CH_3$$

$$CH_3 - C = N$$

$$N = C - CH_3$$

$$N = C - CH_3$$

There are 7 figures, 3 tables, and 11 references, 5 of which are Soviet.

ASSOCIATION: Kishinevskiy gosmdarstvennyy universitet (Kishinev State Univer-

SUBMITTED: June 10, 1957

1. Complex compounds-Spectra 2. Complex compounds-Chemical

reactions 3. Amines-Chemical properties Card 2/2

### "APPROVED FOR RELEASE: 06/13/2000 CIA

CIA-RDP86-00513R000413120004-6

5(2) SOY/78-4-10-6/40 AUTHORS: -Ablov. A. V., Filippov, M. P. Dependence of the Absorption Spectra of the Glyoximines of TITLE: Trivalent Cobalt of the [Co(Amine)(DH) Hal] Type on the Nature of the Co-ordinated Amine PERIODICAL: Zhurnal neorganicheakoy khimii, 1959, Vol 4, Nr 10, pp 2204-2212 (USSR) ABSTRACT: In the formula mentioned in the title D denotes dimethyl glyoxime, hal - chlorine or bromine, amine - ammonia, pyridine, aniline, o-, m-, p-toluidine, o-, m-, p-chloro-aniline or o-, m-, p-bromo-aniline. The production of these non-electrolytes according to the method of L. A. Chugayev (Ref 5) does not yield pure preparations. For this reason the reaction of ammonia and amines with diazido-bis-dimethylglyoxime-cobaltic acids found by A. V. Ablov and N. M. Samus' (Ref 6) was used. The analyses of the preparations are given in table 1. In figures 1-5 the light absorption curves are shown and in table 2 position and intensity of the absorption bands in the region 210-600 mm. In contrast with the dioxime electrolytes of the structure [Co(amine)2(DE)2]+ the compounds investigated Card 1/2

507/78-4-10-6/40 Dependence of the Absorption Spectra of the Clyoximines of Trivalent Cobalt of the [Co(Amine)(DH)2Hal] Type on the Nature of the Co-ordinated Amine

exhibit primary bands in the inner sphere owing to the presence of the halogen, the position of which does not depend on the dipole moment of the aromatic amine, which, however, can be masked by an intense ultraviolet absorption. The band in the range 340-370 mm depends on the nature of the co-ordinated aromatic amine and is shifted into the short-wave range with respect to the band of the dioximine electrolytes owing to the trans-position of the halogen. The band in the range of from 250-300 m/ is due to the Co(DH)2 group. There are 5 figures, 2 tables, and 11 references, 8 of which are Soviet.

Otdel neorganicheskoy khimii Moldavskogo filiala Akademii nauk ASSOCIATION:

SSSR (Department of Inorganic Chemistry of the Moldau Branch

of the Academy of Sciences, USSR)

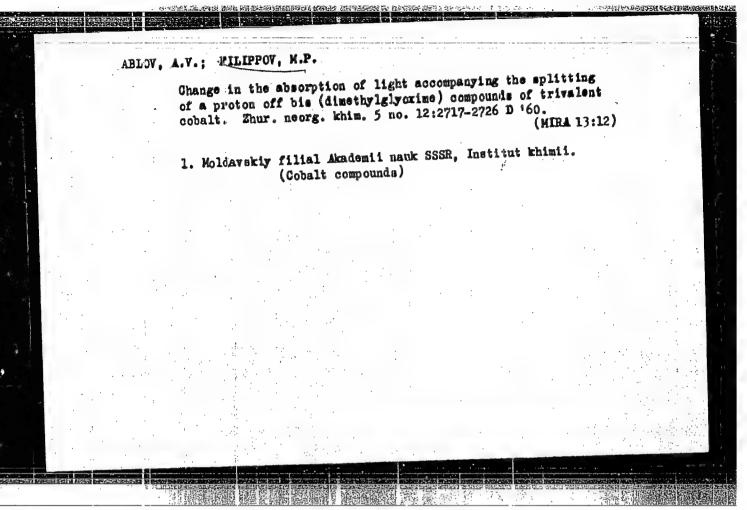
June 19, 1958 SUBMITTED:

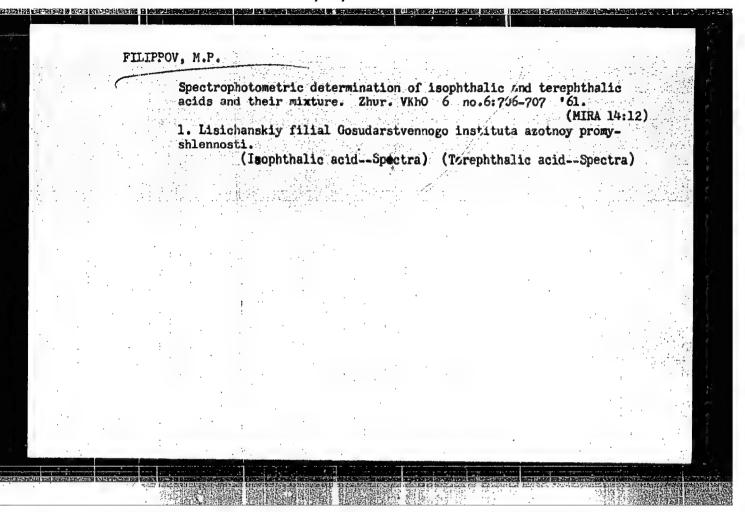
Card 2/2

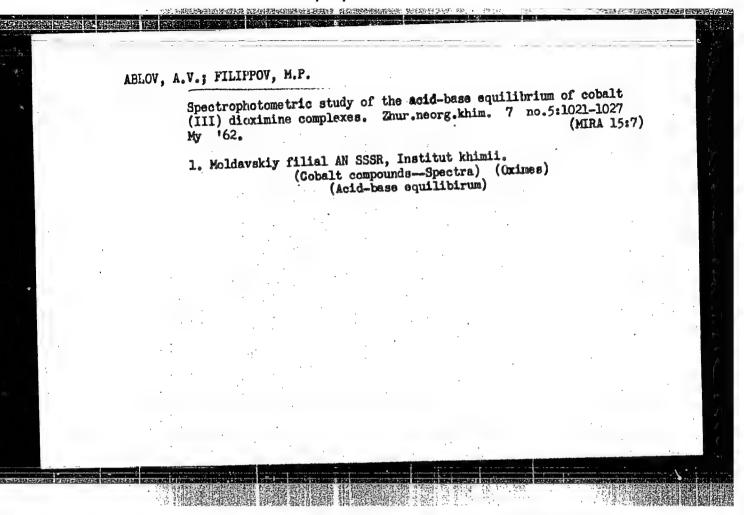
ABLOY, A.V.; FILIPPOV, M.P.; SAMUS', N.N.

Existence of cis- and trans-diaquobis(dimethylglyoximato)
cobaltates(III). Dokl.AN SSSR 133 no.3:575-577 Jl '60.
(MIRA 13:7)

1. Moldavskiy filial Akademii nauk SSSR i Kishinevskiy
gosudarstvennyy universitet. Fredstavleno akad. I.I.Ghernyayevym.
(Cobalt compounds)







FILIPPOV, M.P.

#### PHASE I BOOK EXPLOITATION

SOV/6259

- 'Poltavets, Ivan Mikhaylovich, Faina Fedorovna Sinitsyna, Mark Petrovich Filippov, and Mikhail Panteleymonovich Kolyada
- Ostryye radiatsionnyye porazheniya i ikh lecheniye (Acute Radiation Diseases and Their Treatment) Kiyev, Medgiz UkrSSR, 1962. 154 p. (Series: Biblioteka prakticheskogo vracha) 4180 copies printed.
- Ed.: N. I. Konstantinov; Tech. Ed.: L. A. Zapol'skaya.
- PURPOSE: The book is intended for physicians in all specialities and for students of advanced courses at medical institutes.
- COVERAGE: The book describes methods of treating severe radiation injuries, the treatment of patients with radiation sickness, and the pithological changes occurring in the organism in the course of radiation sickness. Classification, diagnosis, and evacuation of casualties from areas of massive destruction and the organization of dosimetric control among the personnel and

Card 2/5/2

Acute Radiation Diseases and Their Treatment	OV/6259	
in the installations of the civilian defense medical servi are discussed in the light of the most recently promulgate operational procedures. There are 47 references, all Sovietical including three translations.	ce i et,	
TABLE OF CONTENTS:		
Introduction		
Sh. I. Characteniation as -	3	
Ch. I. Characteristics of Injuries Due to Atomic Explosions Unique features of atomic explosions Injurious effects of atomic explosions Damage zones about a center of atomic destruction Characteristics of losses and casualties in atomic explosio Nuclear radiation and nuclear radiation measurement units	5 6 13 ns 16 18	
M. II. Radioactive Substances General characteristics	23	
ard 2/5		

# Spectrophotometric determination of terephthalic acid in a mixture of benzenecarboxylic acids. Zhur.anal.khim. 17 no.5:642-643 Ag (MIRA 16:3)

1. Lisichanskiy filial Gosudarstvennogo nauchno-issledovatel skogo i proyektnogo instituta asotnoy promyshlennosti i produktor organicheskogo simtesa, Severodonatsk.

(Terephthalic acid--Spectra)

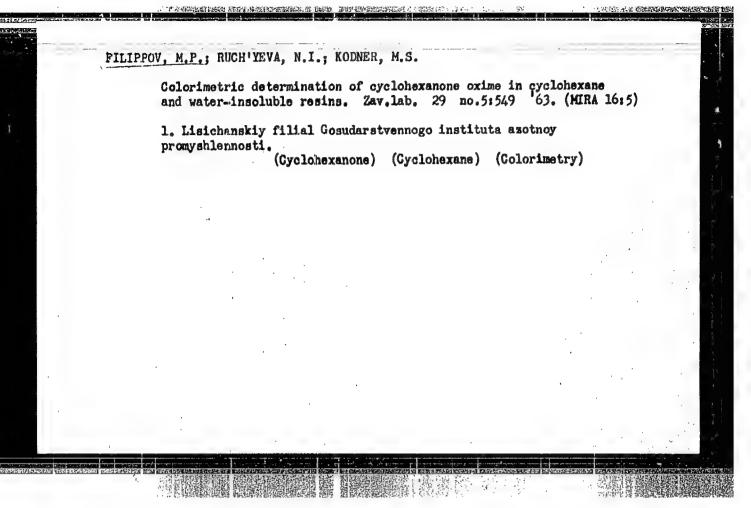
(Benzenecarboxylic acids)

KODNER, M. S.; FILIPPOV, M. P., GUSHCHINA, L. F.

Determination of benzoic, isophthalic, and terephthalic acids in their mixtures. Zhur. VKHO 8 no.2:229-230 163. (MIRA 16:4)

1. Lisichanskiy filial Qosudarstvennogo nauchmo-issledovateliskogo i proyektnogo instituta asotnoy promyshlennosti i produktov organicheskogo sintess.

(Benzoic acid) (Isophthalic acid) (Terephthalic acid)



FILIPPOV, M.P.; VYSOTSKIY, Yu.L.

Cuvette for luminescent analysis at low temperatures. Zav.
lab. 29 no.9:1147-1148 '63. (MIRA 17:1)

1. Lisichanskiy filial Gosudarstvennogo instituta azotnoy promyshlennosti.

RUCE YEVA, N.I.; FILIPPOV, M.P.

Determination of vinyl acetate in its mixture with alkyl vinyl ethers. Zhur. anal. khim. 19 no.3:386-388 '64. (MRA 17:9)

1. Lisichanskiy filial Gosudarstvennogo nauchno-issledovatel'. skogo i proyektnogo instituta azotnoy promyshlennosti i produktov organicheskogo sinteza, Severodonetsk.

FILIPPOV, M.P.; KAGANSKIY, I.M.; PANCHENKO, V.S.; KUTSENKO, V.P.

Spectrophotometric determination of a nitrate ion in complex fertilizers. Zav.lab. 30 no.12:1444-1446 64. (MIRA 18:1)

1. Severodonetskiy filial Gosudarstvennogo instituta azotnoy promyshlennosti.

FILIPPOV, M.P.; ZAYTSEVA, L.F.; ZAYTSEVA, Z.V.; CHUKUR, A.P.

Determination of vinyl alkyl adipates in their mixture with vinyl acetate by the bromide-bromate method. Zhur. anal. khim. 20 no.lx132-134 '65. (MIRA 18:3)

1. Saverodonetskiy filial Gosudarstvennogo nauchno-issledovatel'-skogo i proyektnogo instituta azotnoy promyshlennosti i produktov organicheskogo sinteza.

FILLEPOV, M.P., NUGER, Ta.A.

Formation of molyidenum blue. Zhur. neorg. khim. 16 no.1:
283-285 Ja '65. (MTRA 18:11)

1. Severodonetskiy filial Gosudarstvennogo instituta szetnoy prozyshlennesti. Submitted Febr. 21, 1964.

FILIPPOV, M.P.; IVOIGA, N.F.

Spectrophotometric determination of diphenylolpropane in phenol.
Zhur. VKHO 9 no. 2:234-235 '64. (MIRA 17:9)

1. Lisichanskiy filial Gosudarstvannogo instituta azotnoy promyshlennosti.

#### CIA-RDP86-00513R000413120004-6 "APPROVED FOR RELEASE: 06/13/2000

FILIPPOV, M.S.

AID P - 3519

Sub.ject

: USSR/Power Eng

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Card 1/1

Pub. 26 - 13/30

Authors

Agafonov, M. S., F. T. Makeyev, and M. S. Filippov, Engs.

Title

25 years of operation of the Chelyabinsk State Regional

Power Plant of the Order of Lenin

Periodical

: Elek. sta., 9, 42-43, S 1955

Abstract

The article describes the 25 years of operation of this power plant, without mentioning any engineering details. Names of workers and repairmen are given.

Institution

None

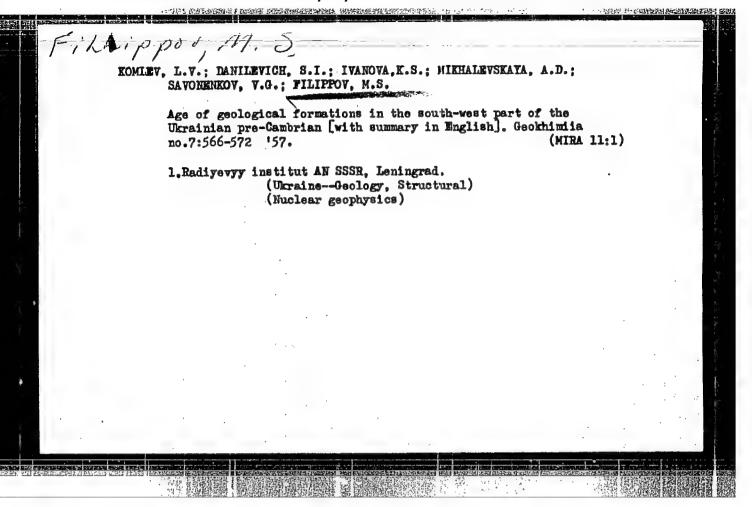
Submitted

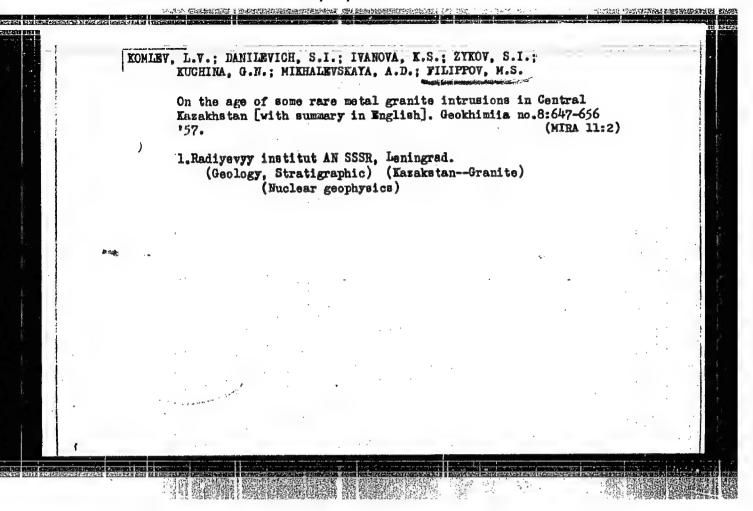
No date

KOMLEV, L.V.; FILIPPOV, M.S.; DANILEVICH, S.I.; IVANOVA, K.S.

Geochemistry of radioactive elements in rocks found in the
Kirovograd - Zhitomir magastic complex in Ukraine. Trudy Radiev.
inst.AN SSSR 7:155-199 '56. (MLRA 10:5)

(Ukraine--Radioactive substances)





FILIPPOV, M. S.: Master Geolog-Mineralo Sci (diss) -- "Radioactive elements in the granites of the central Duepr area". Leningrad, 1958. 23 pp (Leningrad Order of Lenin State U im A. A. Zhdanov), 150 copies (KL, No 6, 1959, 128)

APPROVED FOR RELEASE: 06/13/2000 CIA-RDP86-00513R000413120004-6"

Filippov, M. S.

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FILIPPOV, K. S.

Filippov. M. S. - The Age of Geologic Formations of the South-Mestern Parts of the Ukrainian Pre-Cambrian (Podolia).

The Sixth Session of the Committee for Determining the Absolute Age of Geologic Formations at the Department of Geologic-Geographical Sciences (OGGN) of the USSR Academy of Sciences at Sverdlovsk in May 1957.

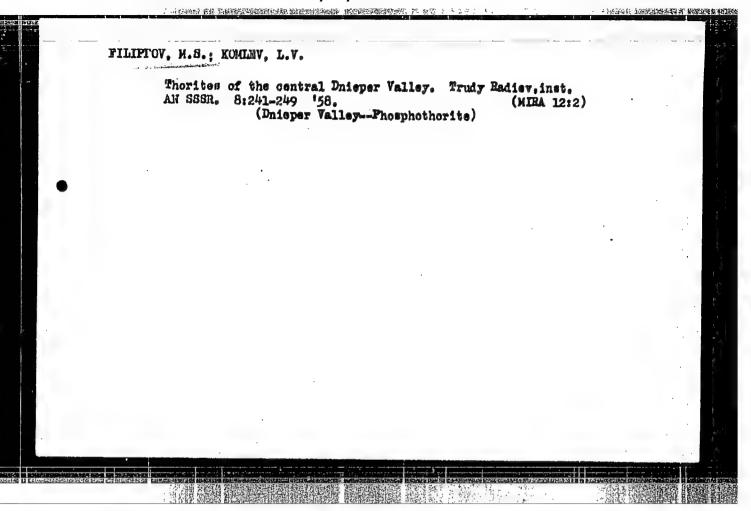
Izv. Ak Mauk SSSR, Ser. Geol., No. 1, 1958, p. 115-117 author Pekarskaya, T. B.

FILIPOV, M.S.

Filippov, M.S., - New Data on the Age of the Ukrainian Pre-Cambrian.

The Sixth Session of the Committee for Determining the Absolute Age of Geologic Formations at the Department of Geologic-Geographical Sciences (OGGN) of the USSR ACademy of Sciences at Sverdlovsk in May 1957

Izv. Ak Nauk SSSR, Ser. Geol., No. 1, 1958, p. 115-117 author Pekarakaya, T. B.



3(8) SOV/7-59-2-3/14

AUTHORS: Komlev, L. V., Filippov, M. S. Danilevich, S. I., Ivanova,

K. S., Kryukova, N. F., Kuchina, G. N., Mikhalevskaya, A.D.

TITLE: Age Data by the Argon and Lead Isotope Method for Some Granites

and Pegmatites of the Central Dnepr Region (Vozrastnyye dannyye argonovogo i svintsovo-izotopnogo metodov dlya neko-

torykh granitov i pegmatitov srednego Pridneprov'ya)

PERIODICAL: Geokhimiya, 1959, Nr 2, pp 110-115 (USSR)

ABSTRACT: This report was presented at the 7th meeting of the Commission for Determination of the Absolute Age of Geological Formations.

An investigation was made of mica from granites and pegmatites, and of accessory monazites and orthites from pegmatite veins. In order to calculate their age from the results of the K/Ar

determination the disintegration constants according to Wetherill et al. were used (Ref 9). For comparative purposes the age was also calculated by the constants found by E. K. Gerling (Ref 10), which had until recently been used in the

Soviet Union for age determinations. Table 1 lists 16 determinations of micas from granites and granodiorites. Values are between 1830 and 2280 million years; biotite from the Yamburg-

Card 1/2 skiy Quarry on the Mokraya Sura River attains 2900 and even

Age Data by the Argon and Lead Isotope Method for Some Granites and Pegmatites of the Central

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2910 million years. Furthermore, two samples each of orthite and monazite were investigated (Tables 2, 3, 4). In order to check the results these analyses were repeated two times. Orthite from Korbino has an age of 2100-2610 million years, biotite from the same place 2280 million years (Table 1). Similarly, it was possible to compare two monazites from the Novo-Danilovskiy Quarry: monazites 1520-2100 million years, biotite 2020 million years. Orthite of Podstepnoye has an age of 2400-3000 million years. This shows that orthite pegmatites may be characterized as relics. There are 4 tables and 12 references, 11 of which are Soviet.

ASSOCIATION: Radiyevyy institut im. V. G. Khlopina, AN ESSR, Leningrad (Radium Institute imeni V. G. Khlopin, AS USSR, Leningrad)

SUBMITTED: July 2, 1958

Card 2/2

THE PROPERTY STATES OF THE PROPERTY STATES OF THE PROPERTY OF SOV/7-59-5-6/14 AUTHORS: Filippov, M. S., Komlev, L. V. TITLE: Uranium and Thorium in the Granitoids of the Middle Pridneprov!ye (Uran i toriy v granitoidakh Srednego Pridneprov'ya) PERIODICAL: Geokhimiya, 1959, Nr 5, pp 437 - 448 (USSR) ABSTRACT: Three complexes of granitoids of the Ukrainian crystalline shield were investigated. The determination of uranium and thorium was carried out in the Laboratoriya geokhimii radioaktivnykh elementov RIAN SSSR (Laboratory of the Geochemistry of the Radioactive Elements RIAN USSR); the activity was measured with electrometers of the type SG-1M. K. S. Ivanova. S. I. Danilevich, V. G. Savonenkov assisted in the investigations. The following complexes were investigated: 1) The oldest complex of granodiorites and plagiogranites has an extraordinarily low content: 1.2.10 $^{-4}$ %U, 0.5.10 $^{-3}$ %Th. With respect to the accessory minerals it belongs to the orthite-sphene-granites. 2) The content of the widely distributed Kirovograd-Zhitomir granites corresponds approximately to the normal Clarke figures (5.7.10-4%), 3.3.10-3%Th). These granites belong, according to the accessory Card 1/2 minerals, to the monazite-garnet group; a part of them to the

Uranium and Thorium in the Granitoids of the Middle SOV/7-59-5-6/14 Pridneprov'ye

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orthite-sphene-granites. 3) The most recent of the three investigated complexes, the Tokovskiy complex, is considerably enriched with thorium and uranium: 9.7.10-3/Th and 9.3.10-4/U. Carrier is above all thorite. The "black quartz" granite of the river Ingulets belongs, according to the accessory minerals to the monazite-garnet group, the tokovskiy granite to the thorite-molybdenit; granites. The results confirm the rule detected by Komlev (Ref 16) that uranium and thorium are enriched in the more recent granites. There are 4 figures, 6 tables, and 16 Soviet references.

ASSOCIATION: Radiyevyy institut im. V. G. Khlopina AN SSSR, Leningrad (Radium Institute imeni V. G. Khlopin AS USSR, Leningrad)

SUBMITTED: October 20, 1958

Card 2/2

### "APPROVED FOR RELEASE: 06/13/2000 CIA-R

CIA-RDP86-00513R000413120004-6

S/015/60/000/009/003/005 A052/A129

AUTHOR:

Filippov. M. S.

TITLE:

Radioactive elements in granites of the central Dnepr region

PERIODICAL:

Referativnyy zhurnal Geologiya, 1960, no. 9, 182, abstract 16996

(Avtoref. diss. kand. geol.-mineralog. n., IGU, Leningrad, 1958)

TEXT: On the basis of three granitoid agglomerates of a different age it is established that the younger rocks have a lower Na, Ca, and Mg content and a higher K, U and Th content compared with more ancient agglomerates. The preferential geochemical combination of U and Th with acid granitoids rich in alkalis is confirmed. [Abstracter's note: Complete translation]

Card 1/1

2006年1月1日 - 1000年1月1日 - 1000年1月 s/081/62/000/012/058/063. B158/B101 Dzhagatspanyan, R. V., Zetkin, V. I., Motsarev, G. V., AUTHORS: Filippov, M. T. Chlorination of silicon-containing monomers and polymers TITLE: under the effect of gamma-radiation Referativnyy zhurnal. Khimiya, no. 12, 1962, 612-613 abstract 12P282 (Sb. "Radioakt. izotopy i yadern. izlucheniya PERIODICAL: v nar.kh-ve SSSR. V. I.". M., Gostoptekhizdat., 1961, 197-200) TEXT: Polydimethylsiloxane rubber (I) and polyphenylmethylsiloxane (II) as well as a number of monomers were chlorinated at 0°C under the action of /-radiation (Co with an activity of 1400 g-equiv of Ra). Chlorination of I takes place easily and rapidly until the introduction of an average of two Cl atoms into the chain of the polymer, after which the process rate falls sharply. In a metal autoclave at both 0°C and 60°C destruction of the polymer takes place. With chlorination of II (molar Card 1/2 36

 Chlorination	of silicon-containing S/081/0	62/000/012/058/063	
atio of Cl:	siloxane = 2:1 and 3:1) substitution a in the aromatic ring. With chlorinat:	and addition chlorination	t
	of Cl <sub>2</sub> :silane = 0.51:1) the basic pro		
erivative;	chlorination of 18.6 g of ethyl-trich	lorosilane (molar ratio	
f Cl <sub>2</sub> :silane	$e = 0.35:1$ ) gives 6.5 g of $\alpha$ and $\beta$ -chi	loroethyl-trichlorosilanes	3 · 4805
$f (c_6H_2cl_3)$	of methyl-phenyl dichlorosilane resul (OCl <sub>3</sub> )SiCl <sub>2</sub> (b. p. 185-188/10 mm). [	lts in the formation	
$f (c_6H_2cl_3)$	of methyl-phenyl dichlorosilane resul (OCl <sub>3</sub> )SiCl <sub>2</sub> (b. p. 185-188/10 mm). [	lts in the formation	190
$f (c_6H_2cl_3)$	of methyl-phenyl dichlorosilane resul (OCl <sub>3</sub> )SiCl <sub>2</sub> (b. p. 185-188/10 mm). [	lts in the formation	30
$f (c_6H_2cl_3)$	of methyl-phenyl dichlorosilane resul (OCl <sub>3</sub> )SiCl <sub>2</sub> (b. p. 185-188/10 mm). [	lts in the formation	30
$f (c_6H_2cl_3)$	of methyl-phenyl dichlorosilane resul (OCl <sub>3</sub> )SiCl <sub>2</sub> (b. p. 185-188/10 mm). [	lts in the formation	90 
hlorination f (C <sub>6</sub> H <sub>2</sub> Cl <sub>3</sub> )( complete tran	of methyl-phenyl dichlorosilane resul (OCl <sub>3</sub> )SiCl <sub>2</sub> (b. p. 185-188/10 mm). [	lts in the formation	30 
$f(c_6H_2cl_3)$	of methyl-phenyl dichlorosilane resul (OCl <sub>3</sub> )SiCl <sub>2</sub> (b. p. 185-188/10 mm). [	lts in the formation	50 84

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8/190/61/003/004/010/014

B101/B207

AUTHORS:

Dzhagatspanyan, R. V., Zetkin, V. I., Motsarev, G. V.,

Filippov. M. T.

TITLE:

Chlorination of organo-silicon monomers and polymers under the action of gamma rays. I. Chlorination of liquid polyphenyl-methyl siloxane and of polydimethyl siloxane rubber.

The infrared spectra of the chlorination products

PERIODICAL:

Vysokomolekulyarnyye soyedineniya, v. 3, no. 4, 1961, 607-612

TEXT: In the introduction, the authors state that initiating the chlorination of organosilicon compounds by means of ultraviolet light proceeds too slowly, however, that chemical initiators as e.g., benzoyl peroxide require a higher temperature at which a sufficient chlorination of methyl chloro silanes is not possible owing to their instability. Therefore, the present study aimed at initiating chlorination by means of gamma rays of Good at low temperatures. The following compounds were chlorinated:
1) Polyphenyl-methyl siloxane (poly-PMS) (molecular weight 2000), and 2) three samples of polymethyl siloxane rubber (poly-MSR) (molecular weight

Card 1/5

21135

S/190/61/003/004/010/014 B101/B207

Chlorination of ...

400,000-500,000). 4-5% solutions of the polymers in CCl $_4$  were used, to which chlorine taken from the cylinder was added. The samples were irradiated in sealed ampoules at 0°C with gamma rays of Co $^{60}$ , activity 1400 g. equ radium. Tables 1 and 2 list the results. Heating with 40% KOH of a chlorinated poly-PMS sample with 55.5% Cl yielded a paste from which crystals with a chlorine content of 64.3-66.7% were separated. On the basis of analytical results, they obtain the empirical formula  $C_6H_6Cl_4$  or  $C_6H_4Cl_4$ . The infrared spectra of the oily residue of hydrolysis showed an intensive

band at 9-10 μ which corresponds to the Si-O bond. Chlorination of poly-MSR led, according to the sample used, to quite different results with respect to the intensity of reaction and the chlorine content of the product obtained. This is due to impurities (catalyst residues) in commercial poly-MSR. Study of the infrared spectra yielded 3690 and 3615 cm<sup>-1</sup> bands both in initial and chlorinated rubber. These bands are due to OH groups (3690 cm<sup>-1</sup> free OH; 3615 cm<sup>-1</sup> OH with H bond). Accordingly, commercial poly-MSR contains silanol groups. As a result of spectral analysis the following is stated: though the IR spectra of chlorinated poly-PMS and poly-MSR differ from those of the initial samples, no absorption bands were found to exist which are characteristic of chlorinated substances.

Card 2, 5

21135

\$/190/61/003/004/010/014 B101/B207

Chlorination of ...

There are 2 figures, 5 tables, and 14 references: 8 Soviet-bloc and 6 non-Soviet-bloc. The 2 references to English-language publications read as follows: Ch. Tamborcki, H. W. Post, J. Org. Chem., 17, 1400, 1952; C. W. Joung, P. C. Servais, C. C. Currie, M. J. Hunter, J. Amer. Chem. Soc., 70, 3759, 1948.

SUBMITTED: July 15, 1960

Onur	3at peni	рушено Гентов, в	Соотно- пление Эмолей Сі осповомоль	Мощность		(G) Bec	(1) Содерінанне жлора, %		
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1 2 3 4 5 6 7	4.76 7.2 4.9 4.9 4.9 2.43 3,3	4,65 4,65 3,12 3,12 3,12 4,08 3,12	1,97:1 2,98:1 3:1 3:1 3:1 1:1 2,03:1	70 70 120 120 120 120 120	30 30 2 5 10 15	8,874 11,425 6,4819 7,0128 7,6840 7,1914 5,9615	48,6 59,5 56,1 54,6 50,7 33,9 50,2	51 61 61 61 61 34,3	

Card 3/5

#### 5/844/62/000/000/066/129 D204/D307

AUTHORS: Dzhagatspanyan, R. V., Zetkin, V. I., Motsarev, G. V.

and Filippov, M. T.

TITLE: The chlorination of phenylmethyldichlorosilane (I) and

dimethyldichlorosilane (II) under the action of pirradi-

ation

Trudy II Vsesoyuznogo soveshchaniya po radiatsionnoy khi-SOURCE: mii. Ed. by L. S. Polak. Moscow, Izd-vo AN SSSR, 1962,

386-389

TEXT: I was chlorinated in sealed ampoules, at 0 and  $20^{\circ}$ C, under & irradiation (~120 r/sec, over 15 or 30 min), with molar ratios (n) of  $\text{Cl}_2$ : I equal to 0.25:1, 0.5:1, and 1:1, since polychlori-

nated silanes are of interest in preparing fluorinated Si-containing monomers and polymers. In contrast to chemically initiated chlorination of I, the present reaction was one of addition of Cl,

into the aromatic ring rather than substitution into the methyl

Card 1/2

The chlorination of ...

\$/844/62/000/000/066/129 D204/D307

group, the main product being a viscous oil, which by chemical and ir spectroscopic tests proved to be  $\mathrm{CH_3} \cdot \mathrm{C_6H_5Cl_6SiCl_2} \cdot \mathrm{A}$  small amount of  $\mathrm{CH_3}$ -chlorinated compounds was also formed. No product in which chlorination of  $\mathrm{CH_3}$ - and  $\mathrm{C_6H_5}$ -groups occurred simultaneously was observed, although it might form in initial mixtures richer in  $\mathrm{Cl_2} \cdot \mathrm{Silane}$  II was similarly chlorinated at  $0^{\circ}\mathrm{C}$ , with n equal to 0.3:1 and 0.5:1, under 2 min doses of f rays at 120 r/sec, to give ~30% yields of the monochloride and 5 to ~17% yields of the dichloride, the latter becoming greater with increasing n. There are 4 tables.

ASSOCIATION:

NII Goskomiteta, Soveta Ministrov SSSR po khimii (NII for Chemistry of the State Committee, Council of Ministers of the USSR)

Card 2/2

S/076/62/036/008/004/011 B101/B144

AUTHORS:

Filippov, M. T., Dzhagatspanyan, R. V., Motsarev, G. V., and

-Zetkin, V. I.

TITLE:

Infrared spectra of organochioussilanes containing chlorine

in the organic group

PERIODICAL: Zhurnal fizicheskoy khimii, v. 36, no. 8, 1962, 1751 - 1754

TEXT: IR spectra of  $GH_3C_6H_5SiCl_2$  (I);  $CH_2ClC_6H_5SiCl_2$  (II);  $CHCl_2C_6H_5SiCl_2$  (III);  $CCl_3C_6H_5SiCl_2$  (IV);  $(CH_3)_2SiCl_2$  (V);  $CH_2ClCH_3SiCl_2$  (VI), and  $CHCl_2CH_3SiCl_2$  (VII) were studied with the following results: (1) The 3.35 and 3.4 $\mu$  bands correspond to the asymmetric and symmetric stretching vibrations of CH in the methyl group. (2) The position of the bands in the range 11-16 $\mu$  strongly depends on the degree of chlorination: The 11.76 - 12.7 $\mu$  band of V in VI becomes weaker and is shifted toward longer waves; in VII it splits into two bands. (3) The 12.58 $\mu$  band of I corresponds to the Si-bound CH<sub>3</sub> group. It changes with the degree of Card 1/2

Infrared spectra of ...

S/076/62/036/008/004/011 B101/B144

chlorination and disappears in IV. (4) The bands of 13-15 $\mu$  for I-IV correspond to the C<sub>6</sub>H<sub>5</sub> groups. (5) The 15.62 $\mu$  band of VI and the 15.38 $\mu$  band of II are ascribed to the SiCH<sub>2</sub>Cl group. There are no bands in this range for the other compounds. (6) In the case of IV, 11.36 and 11.90 $\mu$  bands were observed which appear due to symmetric and asymmetric stretching vibrations of the C-Cl bond in CCl<sub>3</sub>. This was confirmed by the fact that CCl<sub>3</sub>(CH<sub>3</sub>)Si(OC<sub>2</sub>H<sub>5</sub>)<sub>2</sub> and (CCl<sub>3</sub>)<sub>2</sub>Si(OC<sub>2</sub>H<sub>5</sub>)<sub>2</sub> also showed bands in the range 11-11.4 $\mu$  which were absent in compounds containing no CCl<sub>3</sub> group. There are 5 figures and 2 tables.

SUBMITTED: November 9, 1960

Card 2/2

ACCESSION NR 1 AP4034544

S/0020/64/155/005/1163/1166

AUTHORS: Dzhagatspanyan, R.V.; Filippov, M.T.: Motsarev, G.V.; Zetkin, V.I.; Rozenberg, V.R.

TITLE: Radiative chlorination of certain organochlorosilances and

organopolysiloxanes

SOURCE: AN SSSR. Doklady\*, .v. 155, no. 5, 1964, 1163-1166

TOPIC TAGS: chlorination, irradiation chlorination, organochlorosilane, organopolysiloxane, chlorination mechanism, polydimethyl-siloxane, polyphenylmethylsiloxane, ethyltrichlorosilane, methyltrichlorosilane, dimethyldichlorosilane, phenyltrichlorosilane, phenylmethyldichlorosilane, photochemical chlorination, substitution chlorination, addition chlorination, ionic mechanism, free radical

ABSTRACT: The mechanisms involved in the chlorination of various organosilane derivatives under the influence of Co<sup>60</sup> radiation were investigated. A polydimethylsiloxane resin, molecular weight 400,000-500,000, was chlorinated at OC as a 4% solution in CCl<sub>4</sub>. After 'Card-|1/

#### "APPROVED FOR RELEASE: 06/13/2000 C

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ACCESSION NR: AP4034544

chlorination under 4200 rad/min. radiation the chlorine content was 50-55%; optimum reaction time was 15-30 minutes. Total radiation greater than 1.25 x 105 rad did not lead to a higher chlorine content, but promoted degradation of the polymer. By chlorinating polyphenylmethylsiloxane under the same conditions, products containing up to 50.1% chlorine were obtained. About 80% of the chlorine methyl group. Chlorination of ethyltrichlorosilane (molar ratio Cl2: vatives in a ratio of 1:1.7, corresponding to results obtained by and dimethyldichlorosilane the amount of monochloro derivatives in the reaction mixture did not dependon the molar ratio of reagents of chlorination. The relative reaction rate of methyltrichlorosilane for chlorination. The relative reaction rate of methyltrichlorosilane rad/min equaled 0.148 = 0.030 moles/liter-min. The magnitude is proportional to the square root of the power of dosage. The energy

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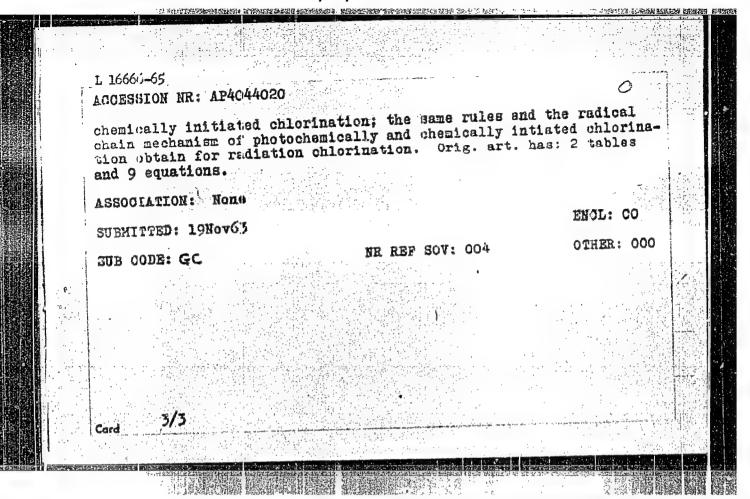
#### CIA-RDP86-00513R000413120004-6

ACCESSION NR: AP4034544 of activation is about 7300-6100 cal/mole for the reaction. Phenyltrichlorosilane and phenylmethyldichlorosilane were chlorinated at 0-150C at 5900 and 800 rad/min at 0-20C. The chlorine added to the double bond of the aromatic nucleus giving C<sub>6</sub>H<sub>5</sub>Cl<sub>6</sub>SiCl<sub>3</sub> and C<sub>6</sub>H<sub>5</sub>Cl<sub>6</sub> (CH<sub>3</sub>)SiCl<sub>2</sub>. This additive chlorination under radiation is analgous to photochemical chlorination. At 500, addition chlorination products as well as products of substition chlorination in the methyl group and the aromatic nucleus were formed. At 100-150C substitution chlorination of the aromatic nucleus predominated indicating ionic mechanism for the arylalkylchlorosilanes. A free radical mechanism was postulated for the alkylchlorosilanes. Orig. art. has: ll equations and 1 table ASSOCIATION: None SUBMITTED: 16Nov63 ENCL: 00 SUB CODE: OC NR REF SOV: 005 OTHER: 002 Card

机自用的复数形式 经股份利用银行 医动物性 医动物性 医动物性 医动物性 医外外外 经工程 计图像 EWT(m)/EPF(c)/EWP(1)/EWA(h)/EWA(1) Pc-li/Pr-li/Fa-i ACCESSION NR: AP4044020 8/0063/64/009/004/0475/0476 AUTHORS: Filippov. M.T.; Dzhagatspanyan, R.V.; Motsarev, G.V.; detkin. TITLE: Radiation thlorination of ethyltrichlorosilane. methyltrichlorosilare and dimethyldichlorosilans SOURCE: Vsesoyuznoye khimicheskoye obshchestvo. Zhurnal, v. 9. no. 4. 1964. 475-476 TOPIC TAGS: radiation chlorination, reaction mechanism, alkylchlorostiant chlorination, liquid phase radiation chlorination, ethylcosilane, methyltrichlorosilane, iimetrv\_ii ..... ane. Systemicalorosilane, chicromethysterials for some and sination immed tion, polychloromethyltrichlorosilane ABSTRICT: The reaction mechanism of radiation-initiated chlorination of alkyl-chlorosilanes was studied. Liquid phase Co-60 radiation-initiated chlorination of ethyltrichlorosilane with molar ratio: of Cl/silane ranging from 0.15 to 0.35 resulted in the forms thon of a and , monochloro derivatives only, with the proportion of  $\beta$ .  $\alpha = 1.72$  when reactant ratio was 0.26 or 0.35, and  $\beta \propto 1$  when Card 1/3

THE PROPERTY OF THE PROPERTY O

L 16665-65 ACCESSION NR: AP4044020 reactant ratio = 0.15 or 0.18. Ohlorination of methyltrichlorsilans with Cl/silane molar ratios ranging from 0.20 to 0.51 gave about 9% .. 171, and a notal concentration of higher only a terivates Listely propertional to the solar catio. Therefore the tose THE INTER STATE OF LOWERS OVER H. . 31 bolycoloro derivatives. Air retarration of the A . server chain mechanism is discussed for the radiation inionination of methyltrichlorosilane wherein the rate of formation of CH2ClSiOl3; and overall reaction is determined by the reaction (4.30), 4 cl-- golding + HCl, and the rate of its disappearance is determined by the reaction OlOHosioly + Ol -- OHOISiOly + HOL. CHoOISiOly is chlorinated about 10 times faster than OH3SiCl3; increasing temperature from 0 to 24.40 increased this chlorination rate about 3 times; time energy of activation is about 7300 cal/mol. The same general rules apply to the chlorination of dimethyldichlorosilane as to methyltrichlorosilane; the rate of the dimethyldichlorosilane chlorination at 00 is 19 times faster than for chlorinating methyltrichlorosilane; its energy of activation is 6100 cal/mol. The effects of the Cl/silane ratio in radiation chlorination are the same as in Card 2/3



MOTSAREV, C.V.; YAKUBOVICH, A.Ya.; ROZENBERG, V.R.; FILIPFOV, M.T.;
DZHAGATSFANYAN, R.V.; BARDENSHTEYN, S.B.; KOLBASOV, V.I.;
ZETKIN, V.I.

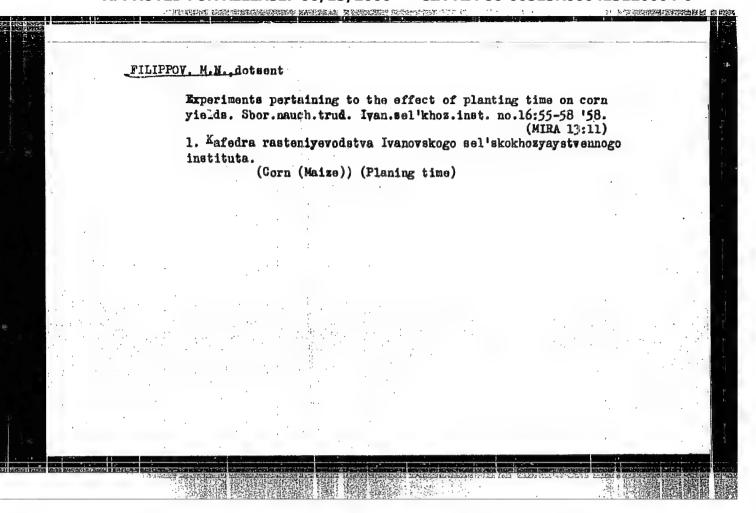
Halogenation of aromatic silanes. Part 17: Addition of chlorine to phenyl-trichlorosilane. Preparation of hexachlorocyclohexyl-trichlorosilane and the mechanism of its formation. Zhur. ob. khim. 35 no.7:1178-1183 J1 '65.

(MIRA 18:8)

SOURCE CODE: VIL/0064/66/000/005/0018/0020 ACC NR: AP6015121 AUTHOR: Dzhagatspanyan, R. V.; Lyaskin, Yu. G.; Filippov, M. T.; Sinitain, V. I.; Yakimenko, L. M.; Glabova, L. I.; Zetkin, V. I. 58 ORG: none TITLE: Radiation chlorination of kerosane SOURCE: Khimichoskaya promyshlennost, no. 5, 1966, 18-20 TOPIC TACS: kerosene, gamma radiation, chlorination, photochemistry ABSTRACT: Groznyy koroseno, from which the aromatic and unsaturated compounds were eliminated by extraction with liquid SO2 was used during chlorination initiated by y-radiation of Co 60 made in the apparatus described by the authors previously (Khim. prom. no. 4, 247, 1965). After purification the kerosene had a molecular weight of 177. Chlorine was passed at the rate of 0.469 g/min in the reactor set into a thermostat with a controlled given temperature. The radiation source was introduced after 15 minutes. The chlorination products were purified from Cl, and HCl by passing a flow of nitrogen. The densities and refractive indexes were measured and the degree of chlorination was determined from the graphs, plotted experimentally, showing the dependence of density de and the refractory indexes not the chlorinated products on their chlorine content. Kinetic curves (content of chlorine vs time in min) were 665.634-4:66.094.403.085.3 1/2 Cord

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L 08659-67 ACC NR: AP6015121 plotted at various temperatures of chlorination (T = 20, 40, and 600) and at various doses of radiation (P = 26.1, 7.3, 1.8, and 0.81 rad/sec). The dependence of the radiation-chemical efficiency coefficient G (number of atoms bound with carbon per 100 equivalent)on the radiation dose P was plotted from kinetic curves. The expression well describes the results obtained. (Dis- $\left(\frac{1600}{7} + 5.76 \cdot 10^{-2} [\% \text{CI}]\right)$ agreement of experimental and calculated values averaged \$\to 10.8%.) This equation can be used for designing a reactor for a temperature range of 0-1000, a radiation dose of 1-50 rad/sec, and a chlorine content of 5-60%. The apparent energy of activation was determined as 3200 cal/mole. The results of radiation chlorination were compared with those of photochemical chlorination and chlorination initiated by azo-bis-isobutyronitryl. It was shown that the same degree of chlorination was achieved more rapidly during radiation chlorination. At T = 200 and P = 26 rad/sec, the product containing Cl>60% was obtained in 90 minutes during radiation chlorination. It took 23 and 21 hours to obtain the same product by photochemical chlorination and chlorination initiated by azo-bis-isobutyronitryl, respectively. Radiation chlorination also has other advantages: it depends little on temperature and is controlled by the radiation dose (easily controllable rate of chlorination), the rate of the radiation process does not depend on the color of the reacting mixture, and there is a much smallor danger of resinification because of an absence of local overheating, Orig. art. has: 3 fig., 4 formulas, and 1 table. SUB CODE: 07/ SUBM DATE: none/ ORIG REF: 001/ OTH REF: 001 Card 2/2



8(3), 8(5) SOY/105-59-8-28/28 RCHTUA Filippov, M. V., Engineer Nomogram for the Determination of the Functions  $\phi(k'h)$  and TITLE: ψ ( k h) PERIODICAL: Elektrichestvo, 1959, Nr 8, Inside of back cover (USSR) ABSTRACT: These two functions serve to determine the resistance of rectangular conductors in the slot of an electric machine. When designing electric machines these functions must be calculated repeatedly for different values of k' and h. This can be done by means of this nomograph. The h-scale is divided into millimeters. An example is given. There is 1 figure. Card 1/1

	THIS I DOOK REPORTED ON \$755	romagnetic Processes in , (Series: Its: Truck	Ed.: A. Tuyleal'bann; Tuch. Ed.: A. Hyerings; Editorial Board: V.O. Tiol, O. T. T. T. Ed.: Editor.; I.M. Kirbo (Resp. Ed.), and Ta. Ta. Klyrial.  Full Editor: I.M. Editor: E intereded for physicists interested in electromagnetic	CONTRACT: This is a collection of fifteen articles by various authors on the investigation of electromagnetic processes in metals by modelling. Indirinal articles treat the fallowing conditions meressary for modelling particular phenomenates are madelling in a variable filed on many modelling the apportantion of ferromagnetic match in a variable filed on an terreted network network of the following the standard of the filed of the ferromagnetic match in a variable filed on	construct resistances; external fields probated by farranges the third have the magnetized in a constant united first orders of the farranges the posts.  Native of united galvante backs and other models for inswitching fields vial continuously distributed abstraction for the probate of inswitching fields fields vial or entire and other resistance in the probate of the arrelated by described in the fields of the probate of the probate of the arrelated probate of the arrelated probate of the probat	with sindian machanical characteristics (revetlantal mentry, period of printtonal, operation with sevenal species of printtonal mentry than the alliest contributions of computing the pensistencial reverse writing on a state of unity; the problems of computing the pensistencial reverse writing on a spiciestes, containing both placed in the trevellate apported field a cytis of difficial and a cytis of a cytis of the cyt	and perfunction of hydromagnetic users of artificate polaritation on the constants of the constants of the constants of the constant of the second of the constant of the constant of the constant flow of lightly dead in indication pergu under the effort of the constants flow of the part of the constants of the constant of th	References accompany the articles.  Estate Lt. Sobiling of the Reletical Field of Electromagnetic Fuge in a Galvanic Bus and on Electrical Comboning Paper	Origorives M.S. Some Problems of Deportiting a System of Interacting Tryliberson's Frictions	Edinia, R.R. Malaticanhip Setmen the Magnetiz Losses in a Furritor Corr With an Open Magnetic Circuit.	Franks, Mark Occidenty Forling or a commercing Administration of States of S	82	Desking, B. Ra., and V. Re. Evertherine. Behavior of Aptronogratio Varua	Kirto, Z.M., Talla, Elyavia, and Z.A. Syntia (Decembed) and L.Ta. Ul'sania. Model of an infinitely form Chammal With Ligaid Serial in a Travelling Magnetic Field.	 Filipper, N.V. Use of Monograms for Rebrinding the Paramters of 15)  Filipper R.V. Nongraphic Calculation of Pasetions  Filipper R.V. Nongraphic Calculation of Pasetions  Filipper R.A. Nongraphic Calculation of Pasetions	Low-Transcruture Induction Benears With 187 Sireniar Gross-Section in the Channel	
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FILIPPOV, M.V.

PHASE I BOOK EXPLOITATION

SOV/6352

Akademiya nauk SSSR. Vychislitel'nyy tsentr

Nomograficheskiy sbornik (Collected Papers on Nomography, no. 1.) Moscow, 1962. 248 p. 1800 copies printed.

Resp. Ed.: G. S. Khovanskiy, Candidate of Technical Sciences; I. A. Orlova; Tech. Ed.: A. I. Korkina.

PURPOSE: This collection of papers is intended for those engaged in research on and design of nomographs.

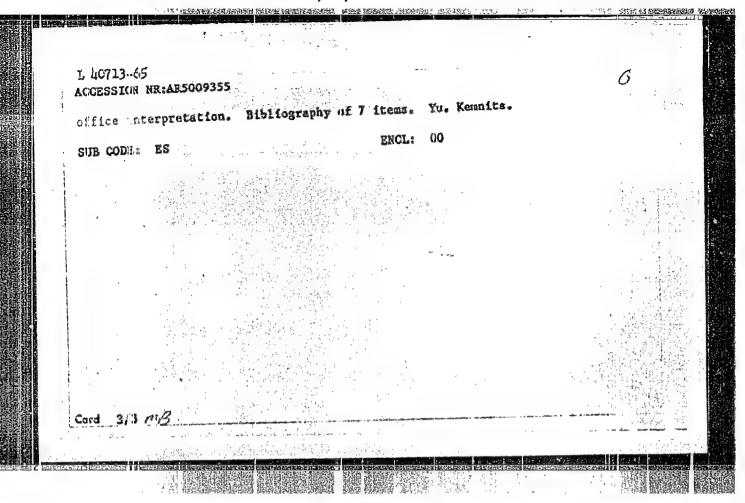
COVERAGE: This collection contains 27 papers concerning various aspects of the theory, construction, and use of nomograms for the solution of algebraic, functional, transcendental, and differential equations. No personalities are mentioned. There are 122 references: 102 Soviet (1 of which is a translation from the English), 8 German, 5 French, 2 English, 2 Spanish, 2 Rumanian, and 1 Czech.

Card 1/10

ollected Papers on Nomography	80V/6352
II. Fel'dman, Ya. S. (Director of the Nomographic Circle at the Leningrad Institute of Precision Mechanics and Optics). The Nomographic Circle of Students in Eligher Technical School	3 1 19
V. Filippov. M. V., Riga. Experience in Using Nomograms in Experimental Investigations	24
. Ul'masov, N., Moscow. Alignment Charts for the Solution of a Transcendental Equation With Three Parameter	s 39
I. Borisov, S. N., Moscow. Constructing Nomograms for a Particular Problem	45
VII. Lapteva, D. G., Moscow. Construction of an Approximate Nomogram by Substituting the Sum of Functions for Their Product	51
VIII. Lapteva, D. G. Construction of a Nomogram with Combined Scales	57
Sard <del>4/10</del> 2/2	

INT(1) AFFTC GW g/0270/65/000/003/0027/0028 0/ ACCESSION NR: AR5009355 SOURCE: Ref. zh. Geodeziya. Otd. vyp., Abs. 3.52,133 AUTHOR: Platonenko, M. A.; Filippov, M. V. Threspretation of agricultural lands on aerial photographs using regression equations while the more representative and the an ibratori da deli bestili CITED : OURCE: Tr. Omskogo s.-kh. in-ta, v.55, no. 2, 1964, 73-80 TOPIC TAGS: serial photography, aerial photograph interpretation, photograpmetry, regression equation TRANSLATION: A statistical study has been made of the influence of the density of the photo tone in the office interpretation of an aerial photograph of agricultural lands. Ten contact prints at a scale of 1:14,000 were used. Between 20 and 30 operact eristic features were noted on each of the prints. The values of the denkirv o the photo tone were determined with an accuracy to 0.25 visual photometric This is an 8-unit scale of a gradation positive. Soils were evaluated with an accours by to 0.5 unit on a 10-unit scale prepared on the basis of the results of **Card** 1/3

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814148 s/057/60/030/009/014/021 B019/B054 and Filippov, M. V. AUTHORS: Characteristics of a Suspended Layer of Ferromagnetic TITLE: Particles in a Magnetic Field Zhurnal tekhnicheskoy fiziki, 1960, Vol. 30, No. 9, PERIODICAL: pp. 1081-1084 TEXT: The suspension and pseudoliquefaction of iron particles (0.1.0.248 cm) in water under the action of an alternating field was carried out with the aid of the experimental arrangement shown in Fig. 1. The particles were placed in a vertical glass tube through which the water was pressed from below. A magnetic coil was arranged around this glass tube, and a small periscope served for the visual observation. It was shown that suspension and pseudoliquefaction of the layer in a magnetic field differ from the same processes in the absence of a magnetic field.

Card 1/2

APPROVED FOR RELEASE: 06/13/2000 CIA-RDP86-00513R000413120004-6"

The authors thoroughly discuss the observations made, and then construct a phase diagram for the state of the suspended layer of ferromagnetic

## 84448

Characteristics of a Suspended Layer of Ferromagnetic Particles in a Magnetic Field

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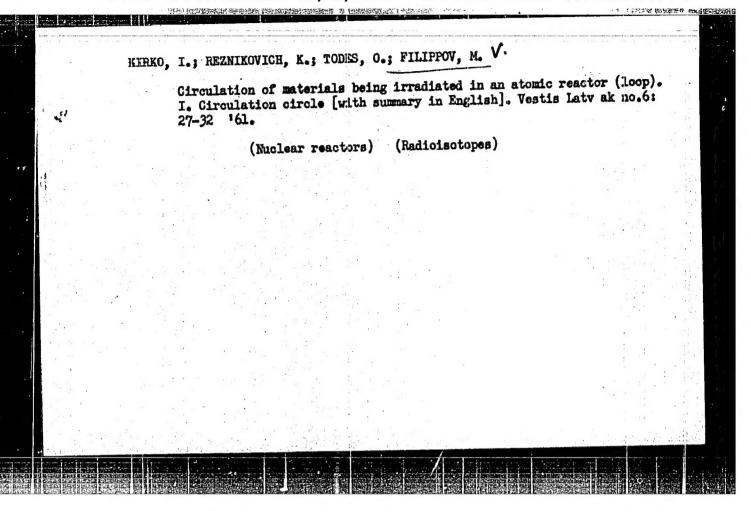
particles in a magnetic field (Fig. 3). This diagram shows on the abscissa the Reynolds numbers, on the ordinate the dimensionless quantity Ma = H<sup>2</sup>h<sub>0</sub>/PD, where h<sub>0</sub> is the initial height of the layer, P its weight, and D the coil diameter. The following phases are shown: layer at rest, pseudopolymeric state, development into pseudoliquefaction, developed pseudoliquid layer, destruction of the layer, and escape of the approximate diagram, in spite of its rough approximation, permits clarifying the rules governing a suspended layer of ferromagnetic particles in a magnetic field. There are 3 figures and 6 references:

ASSOCIATION: Institut fiziki AN Latv. SSR (Institute of Physics of the AS Latviyskaya SSR)

SUBMITTED:

March 31, 1960

Card 2/2



3162h S/197/61/000/012/002/003 B117/B108

14,2110 (1138,1147,1164)

AUTHOR:

Filippov, M.

TITLE:

Effective magnetic permeability of a stratum of ferromagnetic

particles in liquid suspension

PERIODICAL: Akademiya nauk Latviyskoy SSR. Izvestiya, no. 12 (173), 1961,

52 - 54

The effective magnetic permeability  $\mu_{\mbox{\scriptsize eff}}$  of a ferromagnetic stratum is defined by the author as the ratio of the averaged vector of the magnetic induction  $\vec{B}$  to the vector of the external magnetic field strength  $\vec{R}$  in the form of  $\mu_{\text{eff}} = \vec{B}/\vec{H}$ . The directions of  $\vec{B}$  and  $\vec{H}$  do not coincide in general. Such a determination of the effective magnetic permeability is not very accurate, but it has the advantage that the required quantity can be determined by direct measurement. The B component in the direction H was measured for a stratum of magnetite particles of 0.009, 0.012, 0.023 and 0.03 cm diameter, and for a stratum of  $\Phi$ -1000 (F-1000) type ferrite particles of a size of 0.023 cm. The Card 1/3

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Effective magnetic permeability ...

external magnetic field (10 - 100 oersteds). The results were nearly independent of the size of the magnetite particles. The relative height h of the suspension, i. e., the ratio between the height h of the broadening of the stratum in the magnetic field (determined by the demagnetization factor of the "body" = suspension) and the initial stationary height h, was 1.0, 1.25, and 1.5 in all experiments. The effective magnetic permeability of the stratum was found to vary between 2.5, for the smallest fields investigated at the stationary stratum, and ~2.9, at h =1.5 in the strongest field of about 100 oersteds. The effect of the magnetic permeability  $\mu$  of the particles on the quantity  $\mu_{\text{eff}}$  is small, since despite its great diversity for magnetite and ferrite, the corresponding change of  $\mu_{eff}$  does not exceed 20%. The functions  $\mu_{eff} = \mu_{eff}(H)$  show that  $\mu_{\mbox{eff}}$  increases with increasing  $\bar{h}_{\mbox{,}}$  though concentration of the ferromagnetic decreases. The cause for this is the formation of a pseudopolymeric structure in the magnetic field and a decrease in the demagnetization factor of the broadened stratum. In order to clarify the Card 2/3

measurements were made in a homogeneous, longitudinal, variable (50 cps)

3162h S/197/61/000/012/002/003 B117/B108

Effective magnetic permeability ...

effect of the concentration change on  $\mu_{eff}$ , the function  $\mu_{eff} = \mu_{eff}(H)$  was measured at different  $\sigma$  and a constant height of the stratum  $h_o$ . The reduction of  $\sigma$  during the broadening of the stratum is compensated by the orientation and pseudopolymerization of the suspension which involves a decrease in the demagnetization factor. V. G. Vitol is thanked for discussions. There are 4 figures and 3 Soviet references.

ASSOCIATION: In

Institut fiziki AN Latv. SSR (Institute of Physics AS

Latviyskaya SSR)

SUBMITTED:

October 21, 1961

Card 3/3